

REMARKS

Summary of Amendments Made

Claims 1-15 have been amended. Claims 16-31 have been cancelled. Claims 1-15 are now pending. It is believed that no new matter has been added.

The description of the apparatus used in process claims 1 and 3-15 use claim language similar to the process claims of parent serial no. 10/365,300 which in turn is similar to the apparatus claim language from U.S. Patent 6,354,135 which is a parent of this application.

In light of the Quayle action in SN: 10/365,300, a substitute specification and correction to the drawings is also enclosed with this preliminary amendment.

Early and favorable action is earnestly solicited.

Closing

Applicants also believe that this application is in condition for allowance. However, should any issue(s) of a minor nature remain, the Examiner is respectfully requested to telephone the undersigned at telephone number (212) 808-0700 so that the issue(s) might be promptly resolved.

Respectfully submitted,
Norris, McLaughlin & Marcus, P.A.

By: Howard C. Lee
Howard C. Lee
Reg. No. 48,104

220 East 42nd Street
30th Floor
New York, New York 10017
(212) 808-0700

Attachment: Replacement sheet for Figure 1B

CERTIFICATE OF EXPRESS MAILING

I hereby certify that the foregoing Preliminary Amendment (7 pages) is being deposited with the United States Postal Service as an Express Mail, label no. EV 383033753 US to the United States Patent and Trademark Office on the date indicated below:

Date: **29 January 2004**

By: Agata Glinska
Agata Glinska

OBTENTION AND ANALYSIS OF ODORS FROM ODOR EMITTERS

This application is a divisional application of U.S. Serial No. 10/365,300, filed on 12 February 2003, which is a divisional application of U.S. Serial No. 09/999,055, filed on 31 October 2001, now abandoned, which is a divisional application of U.S. Serial No. 09/295,848, filed on 21 April 1999, now U.S. Patent 6,354,135, which claimed priority to Provisional Application No. 60/083,275, filed on 28 April 1998.

FIELD OF THE INVENTION

The present invention relates to obtaining odors from odor emitting objects. More particularly, the present invention relates to an apparatus and process for acquiring odor(s) emitted from odor emitting objects.

BACKGROUND OF THE INVENTION

The emission of odor(s) from objects has given rise to attempts to copy or to acquire odors of interest. For example, aromas from botanical sources, such as living flowers, leaves or other parts of living trees or plants, are sought after in the perfumery arts.

A technique for capturing and analyzing the scent of flowers is described in Perfumes Art Science and Technology edited by P. M. Muller and D. Lamparsky and summarized by R Kaiser in The Scent of Orchids. The method disclosed involves placing a living flower, which is part of a living plant or tree, into an enclosed glass vessel. The glass vessel must be of suitable size and shape to permit the flower to be enclosed without damaging the plant or flower. Specially designed glassware is often required to accommodate particular types of flowers.

When such vessels are employed, the aroma chemicals surrounding a flower. i.e., the headspace, fill the vessel with a vapor phase. The headspace volatiles are drawn through an adsorption trap by means of a pump, over a period of 30 minutes to 2hours. Adsorbents commonly employed in the trap are activated charcoal or special polymeric materials such as TENAX® (2,6-diphenylene polymer) or Porapak Q® (ethylvinylbenzene-divinylbenzene

copolymer). The trapped aroma chemicals are eluted from the trap with a solvent, and injected into a gas chromatograph and analyzed by mass spectrometry (GC/MS).

Mookherjee, et al., U.S. Pat. No. 5,369,978, which is incorporated herein by reference, discloses improvements in the above noted method, although essentially the same principles apply.

In addition, a technique using solid phase micro extraction (SPME) has been described by Mookherjee, et al., Perfumer and Flavorist 23, pp. 1-11 (1998). This technique requires placing a single SPME needle, which is a 2-3 mm solid glass fiber coated with a high boiling point liquid adsorbent, in close proximity to a flower for 30thirty minutes to 60sixty minutes. The aroma molecules are adsorbed onto the needle-like glass fiber and are then analyzed by GC/MS.

Great difficulty, however, has been encountered in attempting to collect scents from certain aroma sources using conventional aroma collecting devices. Special glassware for enclosure of flowers can be inconvenient to use, particularly when the flowers are not readily accessible. Also, they can be awkward to carry to and from remote locations. The rain forest, for example, has a cornucopia of fragrant scent-emitters. However, flowers are often at the end of high slender branches in the forest canopy and are difficult to reach.

Additionally, it is very difficult to attach the conventional collecting devices to the outer branches of trees. Often they do not support the weight of either a person or the equipment employed to collect a sample, particularly in view the length of time required, which can thwart efforts to obtain aromas from some sources.

With respect to the SPME technique, SPME needles are very delicate. The single needle employed to obtain the scent may break at any time in dense canopy. In fact, more often than not, such needles may break during capture in such an environment. Additionally, the time required to adsorb the aroma chemicals using the SPME needles can exceed 30thirty minutes, making it impractical for remote collections.

SUMMARY OF THE INVENTION

The present invention addresses these problems by providing an apparatus for obtaining odor(s) emitted from an odor emitting source. The apparatus of the present invention includes an adsorption unit having at least two openings, which may be one or more capillary tubes. The interior surfaces of the tubes comprise an adsorbent material for adsorbing odor chemicals. A suction device for drawing odor chemicals into the adsorption unit, is positioned for and connected to the tube assembly and a connector connects the tube assembly and the drawing device.

The invention also provides a process for obtaining or capturing odor chemicals. The process includes placing an adsorption unit, which comprises on its interior surface an adsorbent material for trapping aroma chemicals, in proximity to an aroma-emitter, and drawing aroma chemicals into an opening of the adsorption unit with a gas drawing means which is operably connected to the adsorption unit.

Further, the invention allows for the thermal desorption of adsorbed sample, cryogenic focusing, and resolution and identification odor components by GC/MS.

It is an object of the present invention to provide an apparatus for obtaining aroma chemicals from, in particular, difficult to reach aroma emitters.

It also is an object of the present invention to provide an apparatus which is readily portable and is durable for obtaining aroma chemicals from, in particular, a dense canopy.

It also is an object of the present invention to provide a process for obtaining aroma chemicals from, in particular, difficult to reach aroma emitters.

It is a further object of the invention to provide a process for obtaining aroma molecules

which facilitates a short capture time.

DETAILED DESCRIPTION OF THE INVENTION

The apparatus for obtaining odors includes an adsorption unit, a suction device for drawing the odor chemicals into the adsorption unit, and a connector for connecting the adsorption unit and the suction device.

The adsorption unit has an opening for receiving odor chemicals and an opening displaced away from the first for drawing the odor chemicals through. The adsorption unit can be a tube or preferably, a plurality of tubes. A plurality of capillary tubes are the most preferred type of adsorption unit. As used herein, the term odor includes, but is not limited to aromas, as for example, fragrant scents.

The type and dimensions of the tubes of the adsorption unit may vary. For example, a single tube having a plurality of crossing wall portions contained within the tube are considered to be an equivalent alternative to the plurality of capillary tubes. Similarly, tubes having concentric tubes of decreasing diameter contained within a tube also are considered to be equivalent, because of the favorable surface area exposed to odor chemicals. Generally, the inside diameter of the capillary tube ranges from about 0.07 mm to about 1.0 mm and more preferably from about 0.75 mm to about 0.9 mm.

The interior surface(s) of the adsorption unit are coated with an adsorbent material for adsorbing odor chemicals from the air to the stationary material. It is also contemplated that the adsorption unit integrally have interior surface(s) of an adsorbing material. As used herein, an adsorption unit which comprises an adsorbent material on its surface(s) encompasses both of these situations.

The adsorbent employed can vary depending on the odor chemicals sought to be obtained. Generally adsorbent materials, e.g., charcoal, may be used. Preferably, highly adsorbent materials of differing polarity are used to ensure that all aroma chemicals are obtained.

The adsorbent material can be selected from the group consisting of a polar adsorbent, a non-polar adsorbent, an intermediate polarity adsorbent, and any combination thereof. A portion of the capillary tubes may be internally coated with a polar adsorbent, another portion with a non-polar adsorbent, and another portion with an intermediate polarity or neutral adsorbent. Examples of adsorbent materials that are useful in the practice of the invention are Carbowax 20-M as a polar phase adsorbent, methyl silicone as a non-polar adsorbent, and phenyl methyl silicone and polyacrylate as intermediate polarity adsorbents. These materials are coated on the inside of the capillary tube to a thickness of from about 0.1 μ m to about 1.25 μ m, preferably from about 0.1 to about 0.7 μ m, and most preferably from about 0.15 μ m to about 0.7 μ m.

In addition, while preferred tubes for obtaining flower aromas are capillary tubes made from fused silica, alternative tubes may be made of different types of materials which may not be required to be coated per se or which may be made, for example, of metal, of glass, of tempered glass, of a polymer or of a polymer composite.

The length and number of tubes depends upon the scent emitting source. In general, at least one tube is required. Preferably, three tubes of each selected phase are used. The total number of tubes in the bundle is such that the bundle will fit inside a typical thermal desorption tube of a thermal desorption device. For example, the Gertsel Thermal Desorption System TDS 2 instrument requires that the bundle fit into a tube having an internal diameter of about 4 mm. The length of the capillary bundle is from about 5 mm to about 120 mm, usually from about 15 mm to about 120 mm, and more preferably from about 20 mm to about 40 mm.

The capillary bundle is held within the distal end of the connecting tube in an air-tight manner. The connecting tube is of a flexible material such as, for example, PTFE (Teflon), nylon, polypropylene, polyethylene, and the like. The diameter of the connecting tube should be such that the bundle of capillaries fit snugly within an air-tight seal, or another fastening device. Typically, the connection is made of heat-shrink tubing.

The capillary bundle can be suspended in proximity to the aroma emitting source by holding the connecting tube so that the free end of the capillary tubes are held in place, usually over a flower. When the opening of the capillary tubes which is not attached to the connecting tube is placed in proximity with an aroma emitter, the suction device is activated and the aroma chemicals are drawn through the coated capillary tubes from about 2two to about 15fifteen minutes. A 2two to 5five minute collection time is usually sufficient, particularly for the major components. A 5five to 15fifteen minute collection time may be used for components that are present in smaller amounts. The suction device is any device which draws a fluid such as a gas, preferably the headspace air, through each coated capillary tube at a controlled rate. For example, the suction device can be a pump, such as a piston pump, a vane pump, a diaphragm pump, a peristaltic pump or can be a fan. Consequently, a sufficient amount of odor chemicals are adsorbed onto the adsorbent material in a relatively short amount of time to permit analysis of the odor components.

Especially useful are suction devices that do not let the gas being drawn through the capillaries come into direct contact with the suction device itself, thus avoiding contamination. An example of such a suction device is a diaphragm pump that is capable of pumping air from 10-100 ml per minute. The rate can change depending on the composition of the bundle and the number of tubes in the bundle. For example, for a bundle of twenty-four tubes, the optimal flow rate was from about 40 to 60 ml gas per minute. The pump may be operable by conventional means such as by electric current or battery. For trips to remote locations for scent capture, a battery operated pump is used. The pumping rate of the suction device is adjusted using a flow metering device so that the amount of gas passing through the array of capillary tubes in a given time can be measured. Thus, air may be drawn or moved into and through the coated tubes from about 2two to 15fifteen minutes or longer, if desired.

Advantageously, the apparatus has a support for supporting or securing the capillary tubes, and additionally, a housing sheath within which the plurality of capillary tubes are disposed such that they are freely moveable with the connector. Also advantageously, the apparatus includes a means for extending the tube assembly beyond the end of the housing sheath for collection of odor chemicals.

Preferably, the connecting tube holding the capillaries is supported or held by a support such as a rigid rod. The support is of fairly rigid material such as glass, metal, wood or stiff polymeric materials. The support can be any practicable length for reaching odor sources in high branches. Of great use for also reaching aroma sources that are hard to access is a light weight extendible tubular pole which permits the collection device to be extended with the arm portion angled such that the distal end of the device reaches over a flower. The pole can be made of aluminum or the like for a sturdy, light-weight construction. When using an extendible pole, the support can be affixed to the distal end or integral therewith. In addition, the pole can be collapsed for easy transport.

The extendible pole can in practicality be up to 8eight feet long when fully extended, but can reach upwards of 20twenty feet in length, especially when additional sections are connected to the pole. It is advantageous for the extendible pole to be foldable for ease of carrying to about a 3 1/2three and one half feet length. There should be at least one hinging arm portion for extending at a different angle from that of the adjacent portion. If there is only one arm portion, it should be at the distal end for extending the capillaries. How far the pole is extended and at what angle, if any, the arm is extended, depends upon the distance and accessibility of the flower.

Any extension pole used accommodates the connecting tube either alongside or within the pole. To accommodate the connecting tube within the pole, the pole should be hollow, with

enough room in the hollow space of the tubing to run through. There must also be sufficient play for the connecting tube at the hinging points when the arm is angled. It is preferable, however, to run the connecting tube alongside the extendible pole, so that the tube need not be flexed as much by the hinging portion of the arm.

Whether the support at the distal end is integral or attached to the pole, the support itself must support or hold the connecting tube which holds the bundle of capillaries. The support is preferably tubular with a sufficient hollow interior area to accommodate the connecting tube. The support can be attached to the pole such that it can swivel, to provide even further flexibility for accessing a particular flower in dense foliage.

The capillary bundle can be extended to the aroma chemical source by pushing the connecting tube from where the operator stands toward the direction of the aroma chemical source. The connecting tube slides along or within the support and the capillary tubes are thus extended outward. Once the aroma chemicals have been collected, the capillaries are pulled back into the support by pulling the connecting tube. Any manner of moving the connecting tube in the direction of the aroma source is deemed to be within the scope of the present invention.

It is preferable to protect the capillary bundle while it is being extended and maneuvered into position relative to the desired aroma-emitter. We have found that it is desirable to first insert the connecting tube holding the bundle into a housing sheath of semi-rigid material such as nylon which is substantially friction-free with respect to the connector. It is suitable for the housing sheath to be tubular and it is connected or attached to the support or it can be connected or attached within a tubular support. A connecting tube of PTFE (Teflon), for instance, must be freely moveable along the surface of the sheath. Accordingly, the connecting tube holding the capillaries can be extended beyond the end of the housing sheath when in proximity to a flower for collection.

Where the support is tubular and the sheath is attached therein, the support provides the

actual barrier or protective covering to shield the capillary tubes from damage. Yet the sheath which facilitates ready deployment of the capillaries after the apparatus has been extended into position through branches, etc., contributes to the protection of the capillaries.

It is also contemplated that the support can be prefabricated with an inner surface of sheath material.

In the preferred embodiment, the connecting tube within the housing sheath runs alongside the extendible pole. Fasteners can be used at one or more locations along the pole to fasten the sheath to the pole. The housing sheath can be run along the extendible pole beyond any hinge points, or preferably, into the support. This configuration advantageously permits free movement of the connecting tube by the operator pushing the tube at the proximal end of the housing sheath.

When a long pole is used with an attached support at the distal end, the housing sheath can be disposed on or in the support. Preferably, the housing sheath holds the connecting tube within, for the length of the pole. For a hollow extendible pole, the housing sheath runs along its exterior. As the housing sheath readily permits movement of the connecting tube, the connecting tube slides easily forward to extend the tubes for collection when the housing sheath is affixed alongside or throughout the entire length of the pole. It is suitable, though, to affix the housing sheath for about 75% of the length of the pole, where the sheath runs short at the proximal end of the pole.

The capillary bundle containing the trapped aroma chemicals is removed and carefully stored for subsequent processing. The aroma chemicals trapped within the capillaries of a bundle are then placed into a thermal desorber, for example, the Gertsel TDS 2. The desorbed volatiles are then transferred, using an inert gas such as helium at a flow rate of from 5-20 ml per minute, into a cryogenically cooled inlet that uses liquid nitrogen as coolant. The cryogenic focused

materials are then analyzed directly by gas chromatography using an instrument such as a Hewlett Packard 6890 gas chromatograph equipped with a methyl silicone capillary column (0.125 mm x 60 meter), and a mass spectrometer such as the Hewlett Packard 5973 mass spectral detector. Quantitative as well as qualitative analysis of the fragrance components can be performed. The analysis is simplified as the sample can be processed from the desorber through the mass spectrometer by the devices all connected in-line. We have surprisingly found that the process provides data of at least the same quality as the less practical existing methods.

As the present invention permits the capture of the headspace volatiles of a flower high up in the canopy, it will be apparent that when capture is performed in dense canopy, aroma molecules from adjacent aroma-emitters may also be captured. In this case, the captured molecules can represent some combination of aroma molecules from two or more aroma-emitting sources. Analysis of the adsorbed aroma chemicals from more than one source can identify desirable scent combinations.

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1A illustrates a sectional view of a capillary tube bundle held within a connecting tube.

Figure 1B illustrates a cross-sectional view of the capillary tube bundle of Figure 1A.

Figure 2 illustrates the capture of aroma chemicals with a collection apparatus employing an extendible arm and a support attached thereto.

Figure 2a illustrates the support of FIG. 2 in cross-section.

Figure 3 illustrates apparatus for obtaining odors.

Figure 4 illustrates apparatus for analyzing odors.

Figure 5 illustrates a GC graph of resolved chemical components from a 2two minute aroma capture.

Figure 6 illustrates a GC graph as in Figure 5, where aroma capture was for 5five minutes.

Figure 7 illustrates a GC graph as in Figure 5, where aroma capture was for 10ten minutes.

DETAILED DESCRIPTION OF THE DRAWINGS

As illustrated in Figure 1A, a bundle of capillary tubes 10 are held by connecting tube 12. The capillary tubes of bundle of capillary tubes 10 are coated on their inner surfaces (not shown) with adsorbent material(s) of various polarities. Fastener 14 is fastened around connecting tube 12. Figure 1B shows a cross-sectional view of the bundle of capillary tubes 10 from Figure 1A.

Capillary bundle 10 is connected to a pump (not shown in Fig. 1A) by connecting tube 12.

As illustrated in Figure 2, aroma chemicals are obtained with a collection device employing support 26 which is attached by wing nut to extension arm 20. Extension arm 20 represents the distal portion of an extension device. Support 26 can swivel in this embodiment.

Capillary bundle 10 is shown over flower 18 collecting aroma chemicals. Connecting tube 12 is shown extended out from housing sheath 16. Housing sheath 16 runs within support 26 and emerges from aperture 28 of support 26 and runs alongside support 26 and extension arm 20. Fasteners 30 allow for loose fastening of housing sheath 16 to both support 26 and extension arm 20.

Figure 2a2A illustrates a cross-section of support 26. Housing sheath 16 is within support 26, and connecting tube 12 is within housing sheath 16.

Figure 3 illustrates a system for the collection of odors. A collection device is shown with extension device 40 having extension arm 20. Connecting tube 12 runs through housing sheath 16. Housing sheath 16 is within support 26 at its distal end and exits through aperture 28 of support 26 and continues along extension arm 20 and extension device 40. Connecting tube 12 emerges from housing sheath 16 and is operably connected to pump 42.

Figure 4 illustrates the desorption and analysis of the captured aroma chemicals. After capture of aroma chemicals, the capillaries 10 are placed into thermal desorber 44. The desorbed volatiles are transferred into cryogenically cooled inlet 46. The cryogenically focused materials are then analyzed by GC/MS instrument 50.

EXAMPLE 1

Scent was captured using three separate capillary bundles, each bundle having capillaries coated with Carbowax 20M® and capillaries coated with methyl silicone. By use of the aroma collection apparatus and method of the present invention, the aroma of a Hyacinth was collected. For the first capillary bundle, exposure of the coated capillaries to the headspace was for 2two minutes. For the second bundle, the collection time was 5five minutes and for the third, the bundle was exposed for 10ten minutes.

Each bundle of tubes was placed in the thermal desorber. The samples were thermally desorbed, cryogenically focused and then analyzed on a gas chromatography. The gas chromatograph utilized a methyl silicone capillary column (0.125 mm x 60m), a helium gas flow rate of 1.5 ml/min and was programmed from 40°C to 250 °C. at 4° per minute.

Figure 5 is a graphical display of adsorption determined by GC for the first capillary bundle. The intensity of resolved odor chemicals is shown as a function of time. The amounts of the adsorbed components collected over a 2two minute period were sufficient to permit identification of the components by GC/MS.

Figure 6 is a graphical display as in ~~Fig~~Figure 5 for the second capillary bundle. In comparing the relative intensities of corresponding peaks from Figure 5, it can be seen that the intensity of the peaks is greater for a ~~5~~five minute adsorption time than for a ~~2~~two minute adsorption time. All peaks were easily identified by GC/MS.

Figure 7 is also a graphical display as in Figure 5 for the third capillary bundle. In comparing peak intensities, it can be seen that a greater amount of the aroma components can be adsorbed in ~~10~~ten minutes time ~~then~~than for ~~2~~two or ~~5~~five minutes.

As can be seen from Figures 5, 6, and 7, sufficient amounts of aroma chemicals can be collected, in a time period as low as two minutes, in sufficient quantities to permit analysis and identification of those chemicals.

While the invention has been illustrated and described with respect to illustrative embodiment and modes of practice, it will be apparent to those skilled in the art that various modifications and improvements may be made without departing from the scope and spirit of the invention. Accordingly, the invention is not to be limited by the illustrative embodiments and modes of practice.